



## Details for REVA's Annual Meeting

**San Diego, California** (Monday, May 9, 2016, PDT) – As previously announced, REVA Medical, Inc. (ASX: RVA) (“REVA” or the “Company”) will hold its 2016 Annual General Meeting of Stockholders (the “AGM”) on Thursday, May 26, 2016 at 10:30 a.m. in Sydney, Australia (which is 5:30 p.m. on Wednesday, May 25, 2016 PDT). The meeting will be held at the AGL Theatre in the Museum of Sydney located at the corner of Phillip and Bridge Streets.

The AGM will be audiocast and may be accessed toll-free within the United States and Canada by dialing 1-877-312-5413 five minutes prior to the scheduled start time. Callers in Australia may access the call by dialing 1800 005 989. If you are asked to provide an access code, please spell out the word “REVA” to the operator and you will be connected promptly.

If you reside outside of the United States, Canada, or Australia, or if you prefer to access the audiocast through our website, please visit “Events & Presentations” under the “Investors” section of our website at [www.revamedical.com](http://www.revamedical.com), and click on the “listen to webcast” link. A replay of the audiocast will be available on our website after the call.

### About REVA

REVA is a clinical stage medical device company located in San Diego, California, USA, that is working to commercialize its proprietary bioresorbable stents, which are called “scaffolds.” The Company’s scaffolds have been developed as an alternative to metal stents, which are small tube-like devices permanently implanted into an artery to treat coronary artery disease. Scaffolds provide restoration of blood flow, support the artery through the healing process, then disappear (or “resorb”) from the body over a period of time. This resorption allows the return of natural movement and function of the artery, a result not attainable with permanent metal stents. The Company’s initial product, the *Fantom*<sup>®</sup> scaffold, has been designed to offer an ideal balance of thinness and strength and distinct ease-of-use features including complete scaffold visibility under x-ray, expansion with one continuous inflation, and no procedural time limitations. REVA will require successful clinical trial results and regulatory approval before it can commercialize *Fantom* or any other product.

### Forward-Looking Statements

*This announcement contains or may contain forward-looking statements that are based on management's beliefs, assumptions and expectations and on information currently available to management. All statements that are not statements of historical fact, including those statements that address future operating performance and events or developments that we expect or anticipate will occur in the future, are forward-looking statements, such as those statements regarding our ability to obtain regulatory approvals, timely and successfully complete our clinical trials, protect our intellectual property position, commercialize our products if and when approved, develop and commercialize new products, recruit and retain our key personnel, and estimates regarding our capital requirements and financial performance. You should not place undue reliance on forward-looking statements. Although management believes forward-looking statements are reasonable as and*

*when made, forward-looking statements are subject to a number of risks and uncertainties that may cause our actual results to vary materially from those expressed in forward-looking statements, including the risks and uncertainties that are described in the "Risk Factors" section of our Annual Report on Form 10-K filed with the US Securities and Exchange Commission (the "SEC") on March 10, 2016, and as may be updated in our periodic reports thereafter. Any forward-looking statements in this announcement speak only as of the date when made. REVA does not assume any obligation to publicly update or revise any forward-looking statements, whether as a result of new information, future events, or otherwise.*

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